



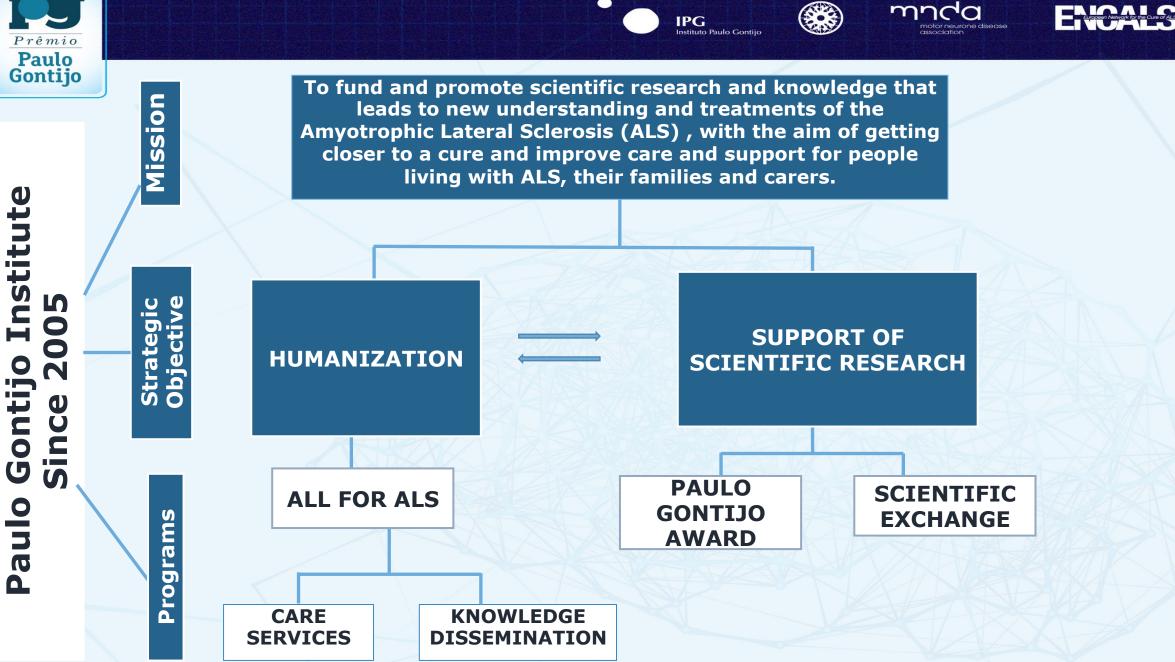
IPG

# Paulo Gontijo Institute- IPG

Brazil

**Dr. Francisco Rotta – Medical coordinator** 















#### FUNDING

- Gontijo Family grant
- Partnerships for our administrative needs: legal, accounting, disclosure, advisory and press
- Listed at the Movement Think&Love site (www.thinkandlove.com.br), a clearinghouse of reputable NGOs that receive donations directly from the site
- Listed at the Paypal donations website https://www.paypal-brasil.com.br/doecompaypal/
- Spontaneous donations usually through PALS and families





- IPG joined in partnership with ABrELA
- Fundraising through social networks, newsletters and emails
- The following campaign was released just as the IBC was reaching Brazilian general public

IPG

Instituto Paulo Gontijo









## PRODUCTION

"Marionette" shows the impact of ALS symptoms and evolution, staring a PALS and a family member

Luciano Szafir, popular Brazilian actor, brother of a PALS Tamara, living with ALS since 2011

#### Cabana Films formed a network of over thirty highly motivated volunteers

It was a technically challenging production, requiring highly skilled professionals

Writer and Director: Lisiana Kieling Executive Producer: Klayane Abreu

XDCAM-HD and XDCAM SD-in 50Mbps TV versions: 15 " and 30 " Theatrical version: 2' Making of version for internet



#### **IPG 2014 Campaign:**

#### Total budget for the Production: US\$ 23,000

## Final cost for the Paulo Gontijo Institute: US\$ 2,500

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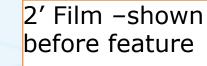
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mnda



 $Pr\hat{e}mio$ 

Paulo Gontijo



European Network for

motor neurone disease

IPG Instituto Paulo Gontijo

6 large metropolitan areas

• Estimated public of 200.000

288 distributed tickets







27,000 new users on the website and social networks

IPG

- 13% returns to the site a high rate of loyalty
- Significant increase in the interactivity of users

IPG raised nearly US\$50,000 that to be used in existing and new programs









# TESTIMONY

**Jocely Pinheiro**: The most important thing is that ALS is starting to become known in Brazil.

#### It is a process!

When my dad was diagnosed in 2012 we did not know almost anything here in Ceará about ALS and now it is already more noticeable thanks to the ants work of patients and families and institutions like this. It is a process in which we are all fundamental participants.











